

FILE 'HCAPLUS' ENTERED AT 15:29:02 ON 01 OCT 2008

L1 183000 S STARCH OR AMYLOPECTIN  
L2 59635 S BRANCHING  
L3 11978 S GELATINIZ?  
L4 47 S L1 AND L2 AND L3  
L5 18 S L4 AND (PY<2000 OR AY<2000 OR PRY<2000)

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```
=> s starch or amylopectin
      181199 STARCH
      7183 AMYLOPECTIN
L1      183000 STARCH OR AMYLOPECTIN
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```
=> s branching
L2      59635 BRANCHING
```

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=> s gelatiniz?
L3      11978 GELATINIZ?
```

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=> s l1 and l2 and l3
L4      47 L1 AND L2 AND L3
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=> s l4 and (PY<2000 or AY<2000 or PRY<2000)
      20096092 PY<2000
      3696782 AY<2000
      3160478 PRY<2000
L5      18 L4 AND (PY<2000 OR AY<2000 OR PRY<2000)
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=> d l5 1-18 ti abs bib
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L5      ANSWER 1 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN
TI      Starch synthase from Canna edulis, its protein and cDNA sequence
        and their use in the production of new starches
AB      The invention provides isolated nucleic acids and their encoded proteins
        that are involved in starch biosynthesis. The invention further
        provides recombinant expression cassettes, host cells, transgenic plants,
        and antibody compns. These nucleic acid mols. can be used to produce
```

transgenic plants having altered structure or quality of starch.  
The present invention provides methods and compns. relating to altering  
the amount and/or morphol. of starch in plants.

AN 2002:551632 HCAPLUS <<LOGINID::20081001>>

DN 137:104816

TI Starch synthase from Canna edulis, its protein and cDNA sequence  
and their use in the production of new starches

IN Singletary, George W.; Zhou, Lan

PA Pioneer Hi-Bred International, Inc., USA

SO U.S., 47 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	US 6423886	B1	20020723	US 1999-388743	19990902 <--
	US 20030135883	A1	20030717	US 2002-44543	20020111 <--
	US 6734341	B2	20040511		
PRAI	US 1999-388743	A	19990902	<--	

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Starch branching enzyme II (SBEII-1 and SBEII-2)  
isoforms from wheat, cDNA, transgenic plants, and altering starch  
properties for food use

AB A class of wheat SBEII genes, SBEII-1, recombinant protein expression in  
transgenic plants, and its use in altering properties of starch  
produced by a plant are claimed. Starch properties include the  
gelatinization onset and/or peak temperature The use of such  
starch with altered properties in food stuff, particularly bakery  
products is also claimed. CDNA clones for SBEII were isolated and  
sequenced. Those clones were divided into two sub-classes, SBEII-1 and  
SBEII-2 having sequence homol. to maize SBEIIb and SBEIIa, resp. These  
genes were mapped to the long arm of wheat group 2 homologous chromosomes.  
Some of those isoforms were expressed as recombinant protein in wheat.  
Differential scanning calorimetry studies showed that starch  
produced in transgenic wheat transformed with expression construct for  
SBEII displayed higher onset, peak, and end temperature for  
gelatinization.

AN 2000:191230 HCAPLUS <<LOGINID::20081001>>

DN 132:247996

TI Starch branching enzyme II (SBEII-1 and SBEII-2)  
isoforms from wheat, cDNA, transgenic plants, and altering starch  
properties for food use

IN Goldsbrough, Andrew; Colliver, Steve

PA Plant Breeding International Cambridge Ltd., UK

SO PCT Int. Appl., 198 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	WO 2000015810	A1	20000323	WO 1999-GB3011	19990909 <--
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW				

RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,  
 ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,  
 CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

AU 9958725	A	20000403	AU 1999-58725	19990909 <--
AU 767103	B2	20031030		
EP 1117814	A1	20010725	EP 1999-946307	19990909 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
HU 2001003618	A2	20020128	HU 2001-3618	19990909 <--
HU 2001003618	A3	20031229		
US 6730825	B1	20040504	US 2001-786480	20010917 <--
US 20040216188	A1	20041028	US 2004-818770	20040406 <--
US 7217857	B2	20070515		
US 20080064864	A1	20080313	US 2007-788837	20070419 <--
PRAI EP 1998-307337	A	19980910	<--	
WO 1999-GB3011	W	19990909	<--	
US 2001-786480	A3	20010917		
US 2004-818770	A3	20040406		

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN  
 TI Molecular background of technological properties of selected starches  
 AB Selected starches, i.e. waxy maize, amaranth, quinoa, wheat, millet, and  
 buckwheat starches, were investigated with respect to their technol.  
 properties such as gelatinization, stability to mech. stress,  
 resistance to conditions, and stability in continuous freeze/thaw cycles.  
 Technol. properties are correlated with mol. features such as  
 branching characteristics in terms of iodine-complexing potential,  
 molar mass, occupied glucan-coil volume, packing d. of glucan coils, and  
 rheol. properties. Waxy maize and amaranth starches were found to be  
 amylopectin-type short-chain branched (scb) glucans with weight average  
 molar masses  $M_w = 17 + 106$  and  $12 + 106$  g/mol, resp. Waxy  
 maize starch had a high gelatinization potential, high  
 viscosity at 95° (340 mPas) low stability at acidic conditions, average  
 stability to shearing, and good freeze/thaw stability. For amaranth  
 starch a viscosity of 122 mPas at 95°, low resistance to  
 acid, but high stability to applied shearing, and even high freeze/thaw  
 stability was determined Investigated quinoa starch was classified  
 as scb-type glucan, however, the branches are significantly longer than  
 those of waxy maize and amaranth. With a  $M_w = 11 + 106$  g/mol and a  
 viscosity of 187 mPas at 95°, this sample is comparably resistant  
 to acidic conditions and to shearing, but instable in freeze/thaw expts.  
 Wheat, millet, and buckwheat starches contain significant percentages of  
 amylose-type long-chain branched (lcb) glucans (22.1, 32.1, and 24.3%,  
 resp.) with  $M_w$  values of  $5 + 106$ ,  $12 + 106$ , and  $15 + 106$   
 g/mol, resp. Wheat starch, with a viscosity of 107 mPas at  
 95°, shows low stability under acidic conditions, but high  
 stability to shearing. Wheat and millet starches, but not buckwheat  
 starch, form weak gels in the course of subsequent freeze/thaw  
 cycles. Millet starch, with a viscosity of 101 mPas at  
 95° was found to be moderately stable under acidic conditions and  
 to shearing. Buckwheat starch with a viscosity of 230 mPas at  
 95° shows no acid resistance and is instable upon shearing but  
 performs very well in freeze/thaw expts.

AN 1999:550347 HCAPLUS <<LOGINID::20081001>>  
 DN 131:171807  
 TI Molecular background of technological properties of selected starches  
 AU Praznik, Werner; Mundigler, Norbert; Kogler, Andreas; Pelzl, Bernhard;  
 Huber, Anton  
 CS Institut Chemie, Univ. Bodenkultur, Vienna, A-1190, Austria

SO Starch/Staerke (1999), 51(6), 197-211  
 CODEN: STARDD; ISSN: 0038-9056  
 PB Wiley-VCH Verlag GmbH  
 DT Journal  
 LA English  
 RE.CNT 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN  
 TI Determination of the distribution of glucose polymers of  
 amylopectin using MALDI-TOF.  
 AB The amount of energy required to gelatinize rice can be crudely  
 determined using the amylose/amylopectin ratio of starch in  
 the rice. However, true energy (cooking time/temperature) requirements are  
 often quite different than predicted values. These discrepancies result  
 in an inconsistent product within the par-boiled rice industry. The  
 differing degree of branching within the amylopectin  
 starch is suspected as the major variable. Current technol. uses  
 gel permeation chromatog. to sep. the debranched chains of  
 amylopectin (glucose polymers) to provide a rudimentary idea of  
 the amylopectin structure. Matrix assisted laser  
 desorption/ionization - Time of Flight Mass Spectrometry (MALDI-TOF)  
 provides a more accurate determination in much less time (45 min vs 5 min).  
 Glucose units with a single Na+ cation attached start at six units and  
 increase in intensity up to 11 units then decrease down to 25 units. It  
 is this distribution of debranched chains that is believed to affect the  
 cooking properties of rice.  
 AN 1999:539290 HCAPLUS <<LOGINID::20081001>>  
 TI Determination of the distribution of glucose polymers of  
 amylopectin using MALDI-TOF.  
 AU Grimm, Deborah A.; Grimm, Casey C.  
 CS Coordinated Instrumentation Facility, Tulane University, New Orleans, LA,  
 70118-5698, USA  
 SO Book of Abstracts, 218th ACS National Meeting, New Orleans, Aug. 22-26 (  
 1999), AGFD-055 Publisher: American Chemical Society, Washington,  
 D. C.  
 CODEN: 67ZJA5  
 DT Conference; Meeting Abstract  
 LA English

L5 ANSWER 5 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN  
 TI A process for textile warp sizing using enzymatically modified starches  
 AB The process comprises the steps of treating a suspension of  
 gelatinized starch with an enzyme selected from the  
 group consisting of cyclodextrin glycosyltransferase, glycosyltransferase  
 and branching enzymes so as to reduce the viscosity of the  
 suspension, and applying the treated starch suspension to  
 textile fibers.  
 AN 1999:451408 HCAPLUS <<LOGINID::20081001>>  
 DN 131:89051  
 TI A process for textile warp sizing using enzymatically modified starches  
 IN Hendriksen, Hanne Vang; Pedersen, Sven; Bisgard-Frantzen, Henrik  
 PA Novo Nordisk A/S, Den.  
 SO PCT Int. Appl., 17 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	WO 9935325	A1	19990715	WO 1998-DK564	19981218 <--

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

AU 9916637 A 19990726 AU 1999-16637 19981218 <--

PRAI DK 1997-1555 A 19971230 <--

WO 1998-DK564 W 19981218 <--

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 6 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Consequences of antisense RNA inhibition of starch branching enzyme activity on properties of potato starch

AB Antisense constructs containing cDNAs for potato starch branching enzyme (SBE) were introduced into potato (*Solanum tuberosum* L.). A population of transgenic plants were generated in which tuber SBE activity was reduced by between 5 and 98% of control values. No significant differences in amylose content or amylopectin branch length profiles of transgenic tuber starches were observed as a function of tuber SBE activity. Starches obtained from low SBE activity plants showed elevated phosphorus content. <sup>31</sup>P-NMR anal. showed that this was due to proportionate increases in both 3- and 6-linked starch phosphates. A consistent alteration in starch gelatinization properties was only observed when the level of SBE activity was reduced to below .apprx.5% of that of control values. Starches from these low SBE activity plants showed increases of up to 5°C in d.s.c. peak temperature and viscosity onset temperature. Studies on melting of crystallites obtained from linear (1 → 4)-α-D-glucan oligomers suggest that an average difference of double helix length of about one glucose residue might be sufficient to account for the observed differences in gelatinization properties. It is postulated that the modification of gelatinization properties at low SBE activities is due to a subtle alteration in amylopectin branch patterns resulting in small changes in double helix lengths within granules.

AN 1998:508745 HCAPLUS <<LOGINID::20081001>>

DN 129:214130

OREF 129:43447a,43450a

TI Consequences of antisense RNA inhibition of starch branching enzyme activity on properties of potato starch

AU Safford, Richard; Jobling, Steve A.; Sidebottom, Chris M.; Westcott, Roger J.; Cooke, David; Tober, Karen J.; Strongitharm, Barbara H.; Russell, Alison L.; Gidley, Michael J.

CS Biosciences Division, Unilever Research, Sharnbrook, MK 441LQ, UK

SO Carbohydrate Polymers (1998), 35(3-4), 155-168

CODEN: CAPOD8; ISSN: 0144-8617

PB Elsevier Science Ltd.

DT Journal

LA English

RE.CNT 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 7 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Manufacture of gelatinized starch liquid with high transparency

AB The title liquid, when incorporated into food-based oils or higher fatty acid alkali salts causing no discoloration and odor due to oxidative

deterioration, is obtained from starch degradation products having >50% fraction with mol. weight range of 20,000-2,500,000, starch degradation products having DE (dextrin equiv) of 1-20, or starch degradation products having cyclic structure and mol. weight of 8000-800,000. Starch degradation products with cyclic structure can be formed by treating a starch compound or mixture with branching enzymes.

AN 1998:42073 HCAPLUS <<LOGINID::20081001>>

DN 128:129399

OREF 128:25397a,25400a

TI Manufacture of gelatinized starch liquid with high transparency

IN Nakamura, Hiroyasu; Hama, Yoshiaki; Okamoto, Harumi; Miyaki, Yasutomo

PA Ezaki Glico Co., Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	JP 10008026	A	19980113	JP 1996-180061	19960619 <--
	JP 3025869	B2	20000327		
PRAI	JP 1996-180061		19960619	<--	

L5 ANSWER 8 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Starch biosynthesis and modification of starch structure in transgenic plants

AB Starch is synthesized through the ADP-glucose pathway, involving the 3 enzymes ADP-glucose pyrophosphorylase, starch synthase, and starch-branching enzyme. ADP-glucose pyrophosphorylase is the key enzyme of the pathway, determining the flux of C into starch. It generates ADP-glucose, which is the substrate for the starch synthases, from glucose-1-phosphate and ATP releasing pyrophosphate. The enzyme is stimulated by 3-phosphoglycerate and inhibited through inorg. phosphate. The starch synthases, which catalyze the transfer of glucose from ADP-glucose to the nonreducing end of a growing  $\alpha$ -1,4-glucan, are divided into 2 classes, the granule-bound starch synthases (GBSS) and the soluble starch synthases (SS). In both classes several isoforms were described from many different plant species. The branching enzyme, which introduces branch points into the amylopectin, can also occur in different isoforms. Other enzymes present in plants, which also act on  $\alpha$ -1,4-glucans, such as the starch phosphorylases, disproportionating enzyme and different starch hydrolases, might also be important for determining the starch structure and, therefore, its processibility. Many aspects of starch synthesis are not fully understood to date. Starch metabolism can be manipulated through genetic engineering, either by the ectopic expression of different heterologous genes, or through the repression of the expression of endogenous genes using antisense RNA technol. This not only allows the functional anal. of starch biosynthetic proteins, but also the manipulation of starch structure in order to widen its industrial applications. In this way many different potato lines were generated, containing either different amts. of starch, or which synthesize a structurally modified starch. These structural changes relate to the amylose content, the phosphate content, or the gelatinization and gelation characteristics of the starch.

AN 1997:568887 HCAPLUS <<LOGINID::20081001>>

DN 127:261734

OREF 127:51129a,51132a

TI Starch biosynthesis and modification of starch structure in transgenic plants

AU Kossmann, J.; Buttcher, V.; Abel, G. J. W.; Duwenig, E.; Emmermann, M.; Froberg, C.; Lloyd, J. R.; Lorberth, R.; Springer, F.; Welsh, T.; Willmitzer, L.

CS Max-Planck-Institut Molekulare Pflanzenphysiologie, Golm, D-14476, Germany

SO Macromolecular Symposia (1997), 120(Functional Polysaccharides II), 29-38

CODEN: MSYMEC; ISSN: 1022-1360

PB Huethig & Wepf

DT Journal

LA English

L5 ANSWER 9 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Physical association of starch biosynthetic enzymes with starch granules of maize endosperm. Granule-associated forms of starch synthase I and starch branching enzyme II

AB Antibodies were used to probe the degree of association of starch biosynthetic enzymes with starch granules isolated from maize (*Zea mays*) endosperm. Graded washings of the starch granule, followed by release of polypeptides by gelatinization in 2% sodium dodecyl sulfate, enables distinction between strongly and loosely adherent proteins. Mild aqueous washing of granules resulted in near-complete solubilization of ADP-glucose pyrophosphorylase, indicating that little, if any, ADP-glucose pyrophosphorylase is granule associated. In contrast, all of the waxy protein plus significant levels of starch synthase I and starch branching enzyme II (BEII) remained granule associated. Stringent washings using protease and detergent demonstrated that the waxy protein, more than 85% of total endosperm starch synthase I protein, and more than 45% of BEII protein were strongly associated with starch granules. Rates of polypeptide accumulation within starch granules remained constant during endosperm development. Soluble and granule-derived forms of BEII yielded identical peptide maps and overlapping tryptic fragments closely aligned with deduced amino acid sequences from BEII cDNA clones. These observations provide direct evidence that BEII exists as both soluble and granule-associated entities. Thus, it is concluded that each of the known starch biosynthetic enzymes in maize endosperm exhibits a differential propensity to associate with, or to become irreversibly entrapped within, the starch granule.

AN 1996:436720 HCAPLUS <<LOGINID::20081001>>

DN 125:81944

OREF 125:15407a,15410a

TI Physical association of starch biosynthetic enzymes with starch granules of maize endosperm. Granule-associated forms of starch synthase I and starch branching enzyme II

AU Mu-Forster, Chen; Huang, Rongmin; Powers, Joseph R.; Harriman, Robert W.; Knight, Mary; Singletary, George W.; Keeling, Peter L.; Wasserman, Bruce P.

CS Dep. Food Sci., Rutgers Univ., New Brunswick, NJ, 08903-0231, USA

SO Plant Physiology (1996), 111(3), 821-829

CODEN: PLPHAY; ISSN: 0032-0889

PB American Society of Plant Physiologists

DT Journal

LA English

L5 ANSWER 10 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Starch for use in paper sizing or coating process



AB The title process used an aqueous size or coating liquid containing converted starch obtained by treating gelatinized starch or a gelatinized modified starch in aqueous medium with a starch-converting enzyme selected from cyclodextrin glycosyl transferases (EC 2.4.1.19) and the branching enzymes (EC 2.4.1.18). The preparation of converted starch in this manner is simpler than that of conventional process and gives retrogradation-resistant starch for good workability.

AN 1996:115231 HCAPLUS <<LOGINID::20081001>>

DN 124:149110

OREF 124:27685a,27688a

TI Starch for use in paper sizing or coating process

IN Bruinenberg, Peter Martin; Hulst, Anne Coenraad; Faber, Ate; Voogd, Roeland Huibert

PA Coooperative Verkoop- en Productievereniging van Aardappelmeel en Derivaten 'AVEBE', Neth.

SO Eur. Pat. Appl., 15 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	EP 690170	A1	19960103	EP 1995-201751	19950627 <--
	EP 690170	B1	20000906		
	EP 690170	B2	20040225		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
	NL 9401090	A	19960201	NL 1994-1090	19940629 <--
	AT 196172	T	20000915	AT 1995-201751	19950627 <--
	ES 2151575	T3	20010101	ES 1995-201751	19950627 <--
PRAI	NL 1994-1090	A	19940629	<--	

L5 ANSWER 11 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Mutant genes at the r and rb loci affect the structure and physico-chemical properties of pea seed starches

AB Mutant genes at two loci, r and rb, known to encode genes affecting the starch biosynthetic pathway, were studied for their effect on the structure and gelatinization of pea seed starches. Comparisons were made using starches from four lines (RRRbRb, rrRbRb, RRrbrb, and rrrbrb), near-isogenic except for genes at these two loci. All the starches had C-type x-ray diffraction patterns, but different contents of 'A' and 'B' polymorphs. The presence of a mutation at either locus increased the 'B' polymorph content in the starches, although the influence of the r mutation was much greater than that of rb. Differences were discovered in the crystalline structure of the rrRbRb starch which correlated with a high content of amorphous phase as well as with the changes in amylopectin structure. In addition, changes in the crystalline structure of this sample correlated with a lack of cooperative transition during starch gelatinization in excess water. The RRrbrb starch had a greatly increased enthalpy of gelatinization in excess water compared with the wild-type starch. It is proposed that this effect is connected with specific charge interactions between the mols. in the starch granule. The rrrbrb starch had parameters of crystalline structure and gelatinization which reflected the different influences of the two genes. With regard to gelatinization, this starch had relatively wide cooperative transition and low enthalpy and a very high peak temperature of transition.

AN 1996:55346 HCAPLUS <<LOGINID::20081001>>

DN 124:85197

OREF 124:16025a,16028a

TI Mutant genes at the r and rb loci affect the structure and physico-chemical properties of pea seed starches  
AU Bogracheva, T. Ya.; Davydova, N. I.; Genin, Ya. V.; Hedley, C. L.  
CS Inst. Biochem. Phys., RAS, Moscow, Russia  
SO Journal of Experimental Botany (1995), 46(293), 1905-13  
CODEN: JEBOA6; ISSN: 0022-0957  
PB Oxford University Press  
DT Journal  
LA English

L5 ANSWER 12 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Modifications of starch during baking: studied through reactivity with amyloglucosidase

AB Conditions that ensure starch hydrolysis by amyloglucosidase in a limited substrate system were worked out. Using these conditions, the degree of access of the enzyme to starch mols. was evaluated in different starch materials. Raw starches of different botanical origins are hydrolyzed at different rates, but starches with limited branching hydrolyze more rapidly. A good example of this is a limit dextrin, which is more susceptible than its parent amylopectin. The effect of gelatinization on the enzymic availability of starch was also studied. It was observed that damaged granules undergo amylolysis much more rapidly than do undamaged ones. Therefore, the extent of amylolysis in a given starch is governed by the degree of granule damage. Starch in bread is hydrolyzed more rapidly and extensively than is that in flour and dough, but no significant differences were found in conventional yeast fermentation between soft and durum wheat. On the other hand, bread obtained by acid fermentation initially undergoes slow amylolysis, although the final level reached is the same as in bread made from the same flour by conventional yeast fermentation

AN 1995:972719 HCAPLUS <<LOGINID::20081001>>

DN 124:28348

OREF 124:5459a,5462a

TI Modifications of starch during baking: studied through reactivity with amyloglucosidase

AU Eynard, Lucia; Guerrieri, Nicoletta; Cerletti, Paolo  
CS Dipartimento di Scienze Molecolari Agroalimentari, Universita di Milano, Milan, I-20133, Italy

SO Cereal Chemistry (1995), 72(6), 594-7

CODEN: CECHAF; ISSN: 0009-0352

PB American Association of Cereal Chemists

DT Journal

LA English

L5 ANSWER 13 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Expression of Escherichia coli glycogen synthase in the tubers of transgenic potatoes (Solanum tuberosum) results in a highly branched starch

AB A chimeric gene containing the patatin promoter and the transit-peptide region of the small-subunit carboxylase gene was utilized to direct expression of Escherichia coli glycogen synthase (glgA) to potato (Solanum tuberosum) tuber amyloplasts. Expression of the glgA gene product in tuber amyloplasts was between 0.007 and 0.028% of total protein in independent potato lines as determined by immunoblot anal. Tubers from four transgenic potato lines were found to have a lowered sp. gr., a 30 to 50% reduction in the percentage of starch, and a decreased amylose/amylopectin ratio. Total soluble sugar content in these selected lines was increased by approx. 80%. Anal. of the starch from these potato lines also indicated a reduced phosphorus content. A very high degree of branching of the amylopectin fraction

was detected by comparison of high and low mol. weight carbohydrate chains after debranching with isoamylase and corresponding HPLC anal. of the products. Brabender viscoamylograph anal. and differential scanning calorimetry of the starches obtained from these transgenic potato lines also indicate a composition and structure much different from typical potato starch. Brabender anal. yielded very low stable paste viscosity values (about 30% of control values), whereas differential scanning calorimetry values indicated reduced enthalpy and gelatinization properties. The above parameters indicate a novel potato starch based on expression of the glgA E. coli gene product in transgenic potato.

AN 1994:319510 HCAPLUS <<LOGINID::20081001>>

DN 120:319510

OREF 120:56089a,56092a

TI Expression of Escherichia coli glycogen synthase in the tubers of transgenic potatoes (Solanum tuberosum) results in a highly branched starch

AU Shewmaker, Christine K.; Boyer, Charles D.; Wiesenborn, Dennis P.; Thompson, Donald B.; Boersig, Michael R.; Oakes, Janette V.; Stalker, David M.

CS Calgene, Inc., Davis, CA, 95616, USA

SO Plant Physiology (1994), 104(4), 1159-66

CODEN: PLPHAY; ISSN: 0032-0889

DT Journal

LA English

L5 ANSWER 14 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Carbon-13 nuclear magnetic resonance studies of chemically modified waxy maize starch, corn syrups, and maltodextrins. Comparisons with potato starch and potato maltodextrins

AB Comparative studies of corn syrups, maltodextrins, chemical modified waxy maize starch, and corn starch were carried out by <sup>13</sup>C NMR techniques. Spectral assignments were made for all materials studied and were checked against independent assignments by proton-C correlation spectroscopy. Degrees of branching and polymerization were estimated for maltodextrins from corn starch and were compared with those of potato maltodextrins in relation to differences in gelling behavior and functionality of corn and potato maltodextrins, resp. Chemical shifts were similar for maltodextrins from corn and potato, as well as wheat amylopectin and amylopectin B. A comparison of solid-state <sup>13</sup>C NMR spectra of corn, wheat, and potato starches reveals their polymorphism, in terms of the number of glucose rings in the unit cell of the amylopectin crystalline regions of starch granules. Gelatinization causes changes in the symmetry of the crystalline regions of amylopectins inside waxy maize starch granules and/or increased mobility of branches in such regions. A broad band in the anomeric region of the solid-state <sup>13</sup>C NMR spectra of waxy maize starch is assigned to the disordered regions of amylopectin in the starch granule structure. Structural details were obtained that are relevant to gelatinization and gelling mechanisms. For corn maltodextrins structural details were obtained concerning the degrees of branching and polymerization, as well as the anomers; such details were significantly different between corn and potato starch maltodextrins.

AN 1991:407164 HCAPLUS <<LOGINID::20081001>>

DN 115:7164

OREF 115:1423a,1426a

TI Carbon-13 nuclear magnetic resonance studies of chemically modified waxy maize starch, corn syrups, and maltodextrins. Comparisons with potato starch and potato maltodextrins

AU Mora-Gutierrez, Adela; Baianu, Ion C.

CS Coll. Agric., Univ. Illinois, Urbana, IL, 61801, USA

SO Journal of Agricultural and Food Chemistry (1991), 39(6),  
1057-62  
CODEN: JAFCAU; ISSN: 0021-8561

DT Journal

LA English

L5 ANSWER 15 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Carbon-13 dioxide breath test to measure the hydrolysis of various  
starch formulations in healthy subjects

AB The 13CO2 starch breath test was used to study the effect of  
physicochem. characteristics of starch digestion. As  
starch is hydrolyzed to glucose, which is subsequently oxidized to  
CO2, differences in 13CO2 excretion after ingestion of different  
starch products must be caused by differences in the hydrolysis  
rate. To study the effect of the degree of chain branching,  
waxy starch, containing 98% amylopectin, was compared with  
high-amylose starch, containing 30% amylopectin, and  
normal crystalline starch, containing 74% amylopectin. The  
effect of the extent of gelatinization was studied by comparing  
extruded starch and crystalline starch. Finally, the  
possible inhibitory effect of adding wheat fiber to extruded  
starch on the hydrolysis rate was studied. The 13CO2 excretion  
for 2-4 h after intake of crystalline starch was significantly lower  
than that of extruded starch. Waxy starch was  
hydrolyzed much faster than was high-amylose starch, but there  
was no significant difference between waxy starch and normal  
crystalline starch. Addition of wheat fiber did not influence the  
hydrolysis rate. The 13CO2 starch breath test is an attractive  
test for the study of factors affecting carbohydrate assimilation.

AN 1990:157113 HCAPLUS <<LOGINID::20081001>>

DN 112:157113

OREF 112:26547a,26550a

TI Carbon-13 dioxide breath test to measure the hydrolysis of various  
starch formulations in healthy subjects

AU Hiele, M.; Ghoo, Y.; Rutgeerts, P.; Vantrappen, G.; De Buyser, K.

CS Gastrointest. Res. Cent., Univ. Hosp. Gasthuisberg, Louvain, B-3000, Belg.

SO Gut (1990), 31(2), 175-8  
CODEN: GUTTAK; ISSN: 0017-5749

DT Journal

LA English

L5 ANSWER 16 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Characterization of starch produced by suspension cell cultures  
of Indica rice (*Oryza sativa* L.)

AB Suspension cultures of rice (*O. sativa*), initiated from seed, produced  
significant amts. of starch. Starch accumulated in  
the cultured cells throughout the growth phase and reached a maximum of 7% of  
the cell dry weight at stationary phase. Starch was present in  
compound granules which were birefringent under polarized light.  
Suspension-culture starch had a higher amylose content and a  
lower gelatinization temperature than rice grain starch.  
Addnl., starch branching enzyme, an enzyme involved in  
starch biosynthesis, was characterized by anion exchange  
chromatog. in culture cells and endosperm. Culture cells had at least 1  
major form of starch branching enzyme which differed  
from the multiple enzyme forms present in endosperm.

AN 1989:21211 HCAPLUS <<LOGINID::20081001>>

DN 110:21211

OREF 110:3565a,3568a

TI Characterization of starch produced by suspension cell cultures  
of Indica rice (*Oryza sativa* L.)

AU Landry, Laurie G.; Smyth, D. A.  
CS Tech. Cent., Gen. Foods Corp., Tarrytown, NY, 10591, USA  
SO Plant Cell, Tissue and Organ Culture (1988), 15(1), 23-32  
CODEN: PTCEDJ; ISSN: 0167-6857  
DT Journal  
LA English

L5 ANSWER 17 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN  
TI Viscosity and gelation characteristics of hydroxyethyl starch  
AB Potato starch (I) had a higher inherent viscosity ( $\eta$ ) than hydroxyethylated I, the  $\eta$  of hydroxyethylated I decreased with increasing SD, and native corn starch (II) had a lower  $\eta$  than I and hydroxyethylated I due to its higher degree of branching. The maximum viscosity and its temperature of I were lower than for hydroxyethylated I, and swelling increased with increasing SD. II gelatinized at higher temps. than hydroxyethylated II, the gelation temperature decreasing with increasing SD. The retrogradation of starch was decreased by etherification, e.g. from 22 to 6% for II.

AN 1982:201554 HCAPLUS <<LOGINID::20081001>>

DN 96:201554

OREF 96:33243a,33246a

TI Viscosity and gelation characteristics of hydroxyethyl starch  
AU El-Hinnawy, S. I.; El-Saied, H. M.; Fahmy, A.; El-Shirbeeney, A. E.; El-Sahy, K. M.

CS Fac. Agric., Ein Shams Univ., Egypt  
SO Starch/Staerke (1982), 34(4), 112-14  
CODEN: STARDD; ISSN: 0038-9056

DT Journal  
LA English

L5 ANSWER 18 OF 18 HCAPLUS COPYRIGHT 2008 ACS on STN

TI Amyloses

AB High mol. weight amylose was produced by gelatinizing a H<sub>2</sub>O suspension of up to 15% starch containing up to 50% amylose, adding to the gelatinized starch at 60° and at pH 5.5 an  $\alpha$ -1,6-glucosidase from the genus Aerobacter, Pseudomonas, Lactobacillus, or Escherichia, cooling the mixture to 45°, and maintaining the mixture at this temperature for 1-2 days to hydrolyze the branched structure of amylopectin into straight chained amylose. Thus, a 5% aqueous suspension of purified amylose starch was heated to 100° with stirring at pH 6.0 and further heated to 130° in a N stream and stirred for 20 min for gelatinization. Thereafter it was quickly cooled to 45° and the pH was adjusted quickly to 4.5; an enzyme of Pseudomonas was added at a concentration of 50 units of enzyme/g of starch. The mixture was maintained at 45° for 1.5 days; thereafter, the precipitated amylose was separated and vacuum dried.

The supernatant was concentrated to half the original volume under vacuum and then kept at 0-5° for 12 hr. The resulting ppts. were separated, washed with warm water, and dried. The yield of the 1st amylose obtained was 40% and that of the amylose from the supernatant 30%, based on the dry raw material. The former amylose has a polymerization degree, as determined with periodic acid, of 780 and the latter amylose a polymerization degree of 130. There was no branching in the former amylose and the latter had on the average 1 branch/mol. Hydrolysis with  $\beta$ -amylase using a 0.5% concentration at pH 6.0 at 55° for 12 hr showed 100% yield of maltose from the former and 85% yield from the latter amylose.

AN 1975:123324 HCAPLUS <<LOGINID::20081001>>

DN 82:123324  
 OREF 82:19727a,19730a  
 TI Amyloses  
 IN Yoshida, Mikihiro; Hirao, Mamoru  
 PA Hayashibara Co., Ltd.  
 SO U.S., 3 pp.  
 CODEN: USXXAM  
 DT Patent  
 LA English  
 FAN.CNT 2

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